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## 18 What is Claimed is:

- A pharmaceutical dosage form for use in a dry powder inhalation devise which comprises:
  - (a) at least one micronized or spray dried solid active ingredient, which active ingredient is soluble in water; and
  - (b) a coating material selected from the group consisting of a fatty acid, an alcohol derivative and a poloxamer, wherein the coating material coats at least partially the surface of the active ingredient.

 The pharmaceutical dosage form as recited in claim 1 wherein the active ingredient has been encapsulated and the coating material partially coats the so-encapsulated active ingredient.

- The pharmaceutical dosage form as recited in claim 1 further comprising a solid, pharmaceutically acceptable carrier excipient and the coating material coats at least partially the surface of the agglomerate or the mixture formed by the active ingredient and the carrier excipient.
- The pharmaceutical dosage form as recited in claim 1 wherein the active ingredient has a mean mass aerodynamic diameter of about 0.5 to about 8 µm.
  - The pharmaceutical dosage form as recited in claim 1 wherein the coating material is a fatty acid sorbitan ester or a PEG ether thereof.
  - 6. The pharmaceutical dosage form as recited in claim 5 wherein the sorbitol derivative is selected from the group consisting of sorbitan mono-cleate, sorbitan trioleate, sorbitan monostearate, sorbitan tristearate, sorbitan monolaurate, sorbitan trinaurate, sorbitan monomyristate, sorbitan trimpristate, sorbitan monopalmitate, sorbitan tripalmitate, PEG sorbitan monopalmitate, PEG sorbitan monostearate, PEG sorbitan tristearate, PEG sorbitan mono-cleate and PEG sorbitan trioleate.